Introduction MR enterography (MRE) aids assessment of small bowel (SB) inflammatory bowel disease (IBD). We aimed to determine the frequency and clinical impact of incidental findings detected by MRE in patients with suspected or known Crohn’s disease (CD).

Methods We conducted a retrospective review of 948 MRE studies performed between June 2009 and December 2012 at our institution. Clinical data (demographics, disease characteristics and therapy) were obtained from electronic patient records. Incidental findings were defined as unexpected lesions in or outside the small intestine, previously unknown or unsuspected at the time of referral and unrelated to IBD.

Results Of 948 MRE studies 445 patients had a diagnosis of IBD, 385 had CD, 54 had ulcerative colitis and 16 had IBD unclassified (IBDU).

Of 385 CD patients, 224 were female, mean age 36 (range 12–72) and median follow up of 4 years (range 0–39). Abnormalities were noted in 285 scans, 162 active non-stricturing, 109 active stricturing and 13-fibrostenosis. Within active groups were 29 fistulae and 12 abscesses in 33 patients. Incidental findings included colitis (10), gallstones (17), ovarian cyst (15), sacroileitis (1), renal cyst (10), hepatic cyst (10), splenic haemangiomata (1), mesenteric abscesses (1), adrenal nodule (2), uterine fibroid (4), chronic pancreatitis (1) and splenomegaly (2) associated with portal vein thrombosis in and varices.

70 studies were performed in UC or IBDU; mean age 34 (range 15–82) 39 were female. Small bowel thickening with signs of active inflammation were seen in 9/13. Other findings included a fluid filled collection in the right ischio-anal fossa, pancreatic divisum, gallstones and liver, ovarian and Nabothian cysts, colitis in 6 and colonic polyps in 1.

Indications for MRE in the non-IBD group (503 patients) included iron deficiency anaemia, abdominal pain, weight loss, diarrhoea, vomiting, abnormal colonoscopy or intra-abdominal abscess. Findings included small bowel thickening (4), sub-acute small bowel obstruction (2), small bowel malignancies (2), small bowel stricture (1) and small bowel intussusception (1). Incidental findings included ovariain, hepatic and renal cysts, adrenal adenoma, ascites, splenic and liver haemangiomata, AAA, PUJ obstruction, liver metastases, gallstones, gallbladder polyph, pelvic abscess, uterine fibroids, large bowel stricture, diverticular disease, cirrhosis, lymphadenopathy, horseshoe kidney, atrophic pancreas and acute appendicitis.

Conclusion A small but significant proportion of patients have important incidental findings at MRE. MRE can add meaningfully to the investigation of SB pathology. A careful selection of patients can be achieved through a collaborative approach between radiologists and clinicians.

Disclosure of Interest None Declared.

Poster presentation III

Colorectal/Anorectal

A NATIONAL SURVEY OF LOCAL HEREDITARY COLORECTAL CANCER SERVICES IN THE UK: A HIGHLY VARIABLE APPROACH?

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Introduction The identification of inherited gastrointestinal disease provides an opportunity to prevent colorectal cancer. Heritable factors contribute about 35% of all colorectal cancer risk which has a significant impact on clinical activity in centres managing colorectal cancer. The British Society of Gastroenterology (BSG) and Association of Coloproctologists of Great Britain and Ireland (ACPGBI), released updated guidelines in 2010 for the management of patients with a family history of colorectal cancer. There is evidence that adherence to these guidelines is highly variable both for endoscopic screening and testing individuals for inherited conditions such as Lynch Syndrome and the Polyposis Syndromes. The aim of this survey was therefore to facilitate understanding of how services for patients with inherited colorectal cancer risk can be improved, and to raise awareness of this issue amongst clinicians.

Methods Following consultation within the BSG Cancer Group, UK Gastroenterologists, Colorectal Surgeons, Clinical and Medical Oncologists were invited to complete a short 10 point questionnaire. This was cascaded by email to 1,798 members of the Royal College of Radiologists (RCR), Association of Cancer Physicians (ACP), the BSG and ACPGBI. We sought their opinion and perception of local hereditary colorectal cancer services, also their adherence to and understanding of current national guidelines.

Results Three hundred and eighty-two members responded to the survey, an overall response rate of 21.3%. Although 69% of respondents felt there was an adequate service for these patients, 64% also believed that another clinician was undertaking this work. There was no apparent patient pathway in 52% of centres, and only 33% maintain a register of these patients. Patients rarely receive initial tumour block testing for Lynch Syndrome. When asked what they would like to augment the service they receive many respondents requested ‘clear guidelines’, ‘pathways’ and dedicated support networks. Many appeared to be unaware of the BSG/ACPGBI guidelines for the management of these patients.

Conclusion There was wide variability in practise and in pathways for hereditary colorectal cancer patients with a perception that they should be managed by another unspecified clinician. BSG/ACPGBI National Guidelines are not adhered to, therefore we recommend improved education, well defined pathways and audit in order to improve care of patients with hereditary colorectal cancer risk.

Disclosure of Interest None Declared.